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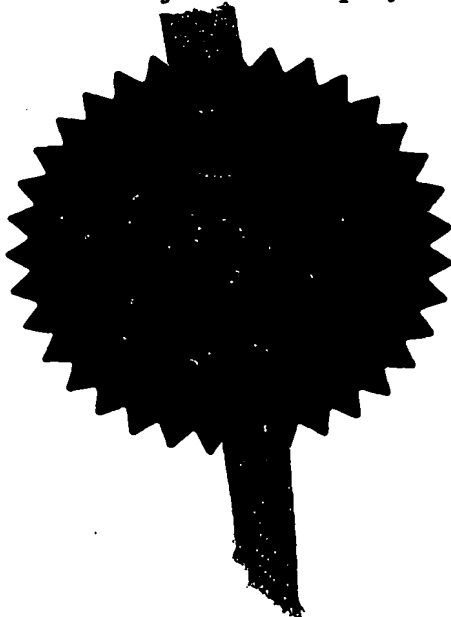
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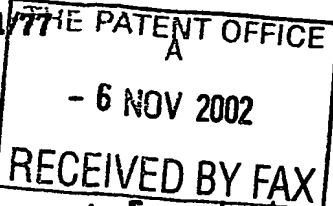
Signed

Stephen Hordley

Dated

24 November 2003

Patents Form 1/77

Patents Act 1977
(Rule 16)06NOV02 176145-1 010763
P01/7700 0.00 0225869.7

Effective Form

The Patent Office

Cardiff Road
Newport
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NP10 8QQ**Request for grant of a patent**

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)

1. Your reference

CDT 370

2. Patent application number

(The Patent Office will fill in this part)

0225869.7

3. Full name, address and postcode of the or of each applicant (underline all surnames)Cambridge Display Technology Limited
Greenwich House
Madingley Rise
Madingley Road
Cambridge CB3 0TX

Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

United Kingdom

6166441004

4. Title of the invention

POLYMER

5. Name of your agent (if you have one)

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

IP Department
Cambridge Display Technology Limited
Greenwich House
Madingley Rise
Madingley Road
Cambridge
CB3 0TX

Patents ADP number (if you know it)

6166441009

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country

Priority application number
(if you know it)Date of filing
(day / month / year)**7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application**

Number of earlier application

Date of filing
(day / month / year)**8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if**

YES

a) any applicant named in part 3 is not an inventor, or
b) there is an inventor who is not named as an applicant, or

c) any named applicant is a corporate body.

See note (d))

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Continuation sheets of this form

Description 19

Claim(s) 4

Abstract -

Drawing(s) -

10. If you are also filing any of the following, state how many against each item.

Priority documents -

Translations of priority documents -

Statement of inventorship and right to grant of a patent (Patents Form 7/77) -

Request for preliminary examination and search (Patents Form 9/77) 1

Request for substantive examination (Patents Form 10/77) 1

Any other documents
(please specify)

11.

I/We request the grant of a patent on the basis of this application.

Signature

Date

08th November 2002

12. Name and daytime telephone number of person to contact in the United Kingdom

Matthew Shade

Tel: 01223 723514

Warning

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent for the same invention and either no direction prohibiting publication or communication has been given, or any such direction has been revoked.

Notes

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- If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.
- If you have answered 'Yes' Patents Form 7/77 will need to be filed.
- Once you have filled in the form you must remember to sign and date it.
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[DUPLICATE]

Polymer

Field of the Invention

This invention relates to semiconductive polymers, their synthesis and use in optical devices.

Background of the Invention

Electroactive polymers are now frequently used in a number of optical devices such as in polymeric light emitting diodes ("PLEDs") as disclosed in WO 90/13148, photovoltaic devices as disclosed in WO 96/16449 and photodetectors as disclosed in US 5523555.

A typical PLED comprises an organic electroluminescent layer located between an anode and a cathode. In operation, holes are injected into the device through the anode and electrons are injected into the device through the cathode. Holes enter the highest occupied molecular orbital ("HOMO") of the electroluminescent polymer and electrons enter the lowest unoccupied molecular orbital ("LUMO") and then combine to form an exciton which undergoes radiative decay to give light. The colour of light emitted from the electroluminescent polymer depends on its HOMO-LUMO bandgap.

An electron transport material is commonly used to assist in transport of electrons from the cathode to the LUMO of the electroluminescent polymer and thus increase device efficiency. Suitable electron transport materials are those having a LUMO level falling between the LUMO level of the electroluminescent polymer and the workfunction of the cathode. Similarly, a hole transporting material having a HOMO level falling between the workfunction of the anode and the HOMO level of the emissive material is commonly used. For example, WO 99/48160 discloses a blend of a hole transporting polymer, an electron transporting polymer and an electroluminescent polymer. Alternatively, the electron transporting functionality and the emissive functionality may be provided by different blocks of a block copolymer as disclosed in WO 00/55927.

A focus in the field of PLEDs has been the development of full colour displays for which red, green and blue electroluminescent polymers are required – see for example Synthetic Metals 111-112 (2000), 125-128. To this end, a large body of work has been reported in the development of electroluminescent polymers for each of these three

colours with red, green and blue emission as defined by PAL standard 1931 CIE co-ordinates.

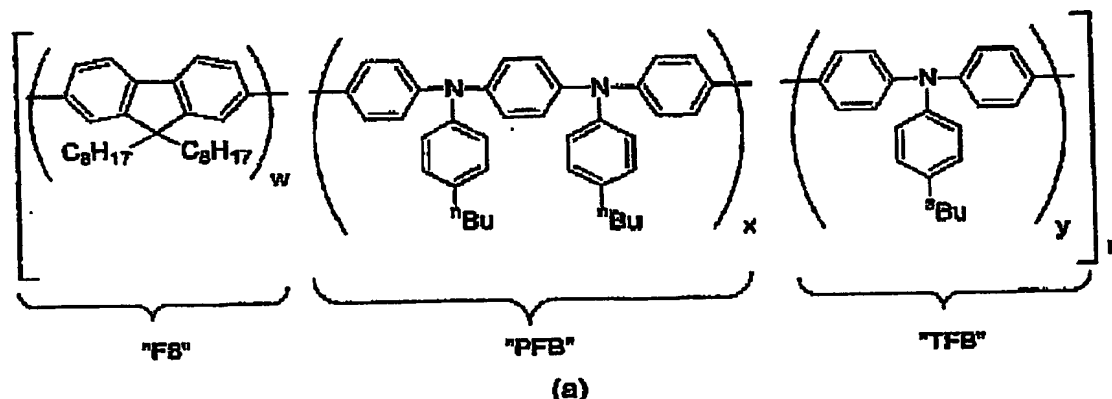
A difficulty encountered with blue electroluminescent polymers is that their lifetime (i.e. the time taken for brightness to halve from a given starting brightness at fixed current) tends to be shorter than that of corresponding red or green materials. One of the factors that has been proposed as contributing to the more rapid degradation of blue materials is that their LUMO levels, and consequently the energy level of the charged state following injection of an electron into the LUMO, tends to be less deep than those of corresponding red or green materials. It is therefore possible that materials comprising these lower electron affinities are less electrochemically stable and therefore more prone to degradation.

For simplicity, a full colour display will preferably have the same cathode material for all three electroluminescent materials. This results in the further problem that the energy gap between the LUMO and the workfunction of the cathode for a typical blue electroluminescent polymer is greater than that for a typical red or green electroluminescent polymer. This may contribute to lower efficiency.

Clearly, assisted electron injection into blue electroluminescent polymers is desirable, however the choice of electron transporting material is constrained by the fact that the emissive material is generally that with the smallest bandgap. This limitation is particularly restrictive in the case of blue electroluminescent polymers since the bandgap required for blue emission is the largest of red, green and blue.

Chains of fluorene repeat units, such as homopolymers or block copolymers comprising dialkylfluorene repeat units, may be used as electron transporting materials. In addition to their electron transporting properties, polyfluorenes have the advantages of being soluble in conventional organic solvents and have good film forming properties. Furthermore, fluorene monomers are amenable to Yamamoto polymerisation or Suzuki polymerisation which enables a high degree of control over the regioregularity of the resultant polymer.

One example of a polyfluorene based polymer is a blue electroluminescent polymer of formula (a) disclosed in WO 00/55927:

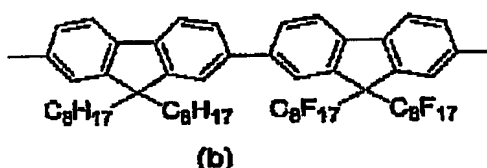


wherein $w + x + y = 1$, $w \geq 0.5$, $0 \leq x + y \leq 0.5$ and $n \geq 2$.

In this polymer, chains of dioctylfluorene, denoted as F8, function as the electron transport material; the triphenylamine denoted as TFB functions as the hole transport material and the bis(diphenylamino)benzene derivative denoted as PFB functions as the emissive material.

WO 94/29883 discloses use of electron withdrawing groups, particularly nitrile groups, as substituents on electroluminescent polymers for the purpose of reducing the barrier to electron injection between a high workfunction electrode and the electroluminescent polymer. This document only teaches use of such substituents on poly(arylene vinylenes).

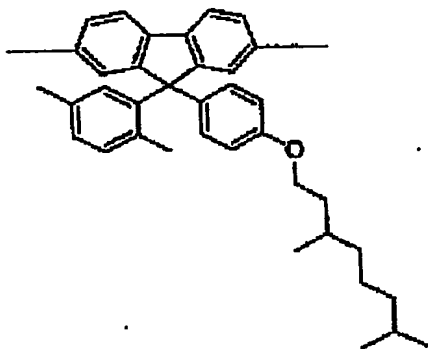
J. Poly. Sci. Part A: Polym. Chem. Vol. 39 (2001) discloses a polymer of repeat units of formula (b):



This disclosure describes use of fluorinated sidechains as a means of decreasing interchain interactions that have been reported to cause aggregation of polyfluorenes

and contains no discussion of using such electron deficient substituents as a means to increase electron affinity. This polymer is disclosed as showing no photoluminescence.

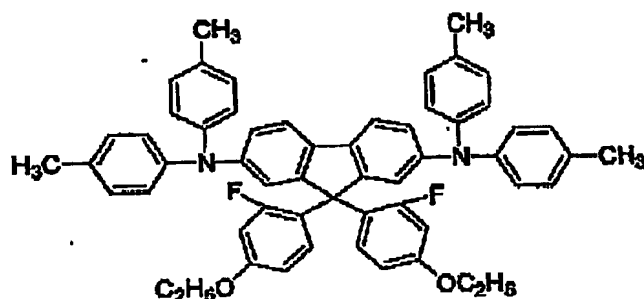
There are disclosures of diphenylfluorenes wherein the phenyl group carries substituents, however these substituents are electron donating as measured by their Hammett sigma constants. For example, WO 00/22026 discloses a homopolymer having a repeat unit of formula (c):



(c)

Also disclosed in this document are copolymers of (c) with dialkylfluorene repeat units and with triarylamine repeat units. Asymmetric substitution of the 9-position of fluorene is described for the purpose of avoiding polymer aggregation; this document contains no teaching of 9-substituents used for the purpose of enhanced electron injection of the fluorene backbone. Similarly, WO 99/20675 discloses a 1:1 copolymer of 9,9-di-n-octylfluorene and 9,9-di(4-methoxyphenyl)fluorene and WO 01/62822 discloses a polyfluorene with triarylamine 9-substituents.

JP 10095972 discloses a molecule of formula (e):



(e)

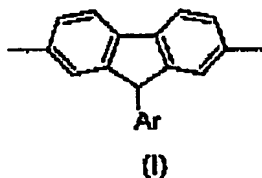
This is disclosed as an emissive material of the type known as "small molecules" rather than polymers as described hereinbefore. This molecule is used in conjunction with a separate, electron transporting molecule. The use of fluorine substituents on the phenyl ring is not described for the purpose of increasing electron affinity of the fluorene ring; fluorine substituents are merely one of a large number of possible substituents for the phenyl ring disclosed in this document.

It is an object of the invention to provide a high electron affinity material that is capable of functioning as an electron transport material for a blue electroluminescent material. For the reasons explained above, such a material would also be capable of functioning as an electron transport material for a red or green material. Furthermore, such material may, as a result of its large HOMO-LUMO bandgap, be used as a blue electroluminescent material.

Summary of the Invention

The present inventors have determined that improved electron injection, and therefore improved PLED performance, may be accomplished by increasing the electron affinity of known polyfluorenes.

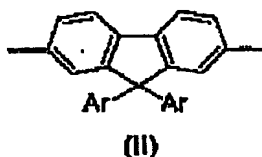
Accordingly, in a first aspect the invention provides a polymer comprising optionally substituted first repeat units of formula (I):



wherein Ar is selected from:

- (a) aromatic hydrocarbon substituted with at least one electron withdrawing group or
- (b) electron withdrawing heteroaryl.

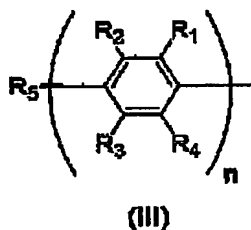
Preferably, the polymer comprises repeat units of formula (II):



wherein each Ar is independently selected from:

- (a) aromatic hydrocarbon substituted with at least one electron withdrawing group or
- (b) electron withdrawing heteroaryl.

Preferred Ar groups according to (a) are independently selected from units of formula (III):



wherein n is from 1-3 and R₁-R₅ are independently selected from:

- hydrogen;

- solubilising groups selected from alkyl, alkoxy, arylalkyl and heteroarylalkyl; and
- electron withdrawing groups

such that at least one of R_1 - R_5 is an electron withdrawing group. Most preferably $n = 1$, i.e. Ar is phenyl.

Another preferred set of Ar groups according to (a) are fused aromatic hydrocarbons such as naphthalene and anthracene.

Preferably, the electron withdrawing group is selected from: groups comprising fluorine, cyano, nitro, carboxyl, amides, ketones, phosphinoyl, phosphonates, sulfones and esters. More preferably, the at least one electron withdrawing group is selected from fluorine atoms, fluoroalkyl, fluoroaryl and fluoroheteroaryl.

Preferred electron withdrawing heteroaryls according to (b) are optionally substituted N-containing heteroaryls, in particular optionally substituted pyridines, most particularly pyridine-4-yl; pyrazines; pyrimidines; pyridazines; triazines, most particularly 1,3,5-triazine-2-yl and oxadiazoles. The electron withdrawing heteroaryl may be substituted with electron withdrawing groups as outlined above to further increase its electron withdrawing effect.

Preferably, the polymer according to the invention comprises a second repeat unit. More preferably the second repeat unit is selected from triaryl amines and heteroaromatics.

Preferably, the polymer according to the invention is capable of transporting electrons. In addition, the polymer preferably has at least one segment capable of hole transport and / or emission. Two or more functions of hole transport, electron transport and emission may be provided by the same segment. In particular, a single segment may function as both an electron transporter and an emitter.

In a second aspect, the invention provides an optical device, preferably an electroluminescent device, comprising a polymer as described above.

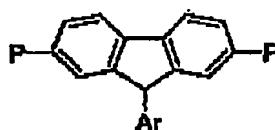
In one embodiment of the second aspect is provided an electroluminescent device comprising:

- a first electrode for injecting charge carriers of a first type;

- a second electrode for injecting charge carriers of a second type; and
- an emissive layer comprising a polymer according to the first aspect of the invention between the first and second electrodes.

The emissive material within the emissive layer may be the polymer according to the first aspect of the invention or another material, preferably another polymer, blended with the polymer according to the first aspect of the invention. Preferably, the polymer according to the first aspect of the invention is capable of transporting electrons in this device.

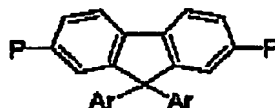
In a third aspect, the invention provides a monomer comprising an optionally substituted compound of formula (IV):



(IV)

wherein each P independently represents a polymerisable group and Ar is as defined above.

Preferably, the monomer comprises an optionally substituted compound of formula (V):



(V)

Preferably, each P is independently selected from a reactive boron derivative group selected from a boronic acid group, a boronic ester group and a borane group and a reactive halide group.

In a fourth aspect, the invention provides a process for preparing a polymer comprising a step of reacting a first monomer as described above with a second monomer that may be the same or different from the first monomer under conditions so as to polymerise the monomers.

Preferably, the process comprises polymerising in a reaction mixture:

- (a) a monomer according to the third aspect of the invention wherein each P is a boron derivative functional group selected from a boronic acid group, a boronic ester group and a borane group, and an aromatic monomer having at least two reactive halide functional groups; or
- (b) a monomer according to the third aspect of the invention wherein each P is a reactive halide functional group, and an aromatic monomer having at least two boron derivative functional groups selected from boronic acid groups, boronic ester groups and borane groups; or
- (c) a monomer according to the third aspect of the invention wherein one P is a reactive halide functional group and one P is a boron derivative functional group selected from a boronic acid group, a boronic ester group and a borane group,

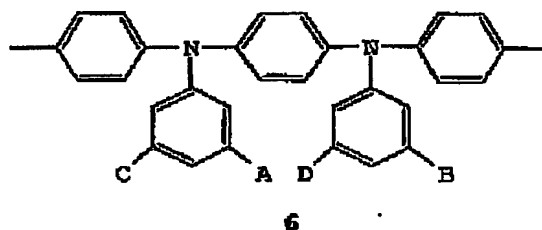
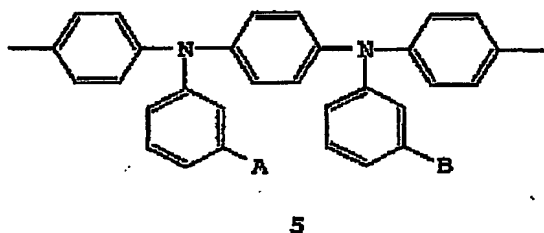
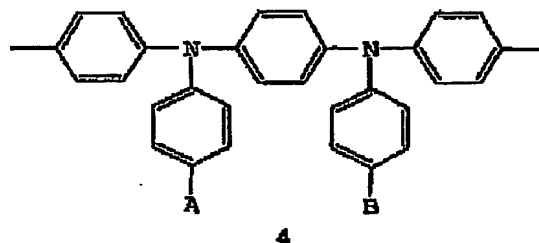
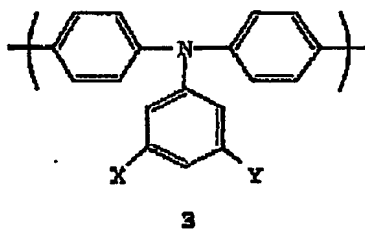
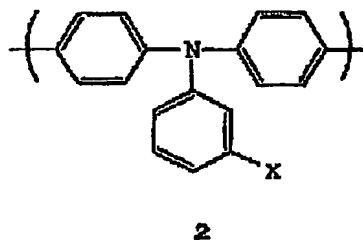
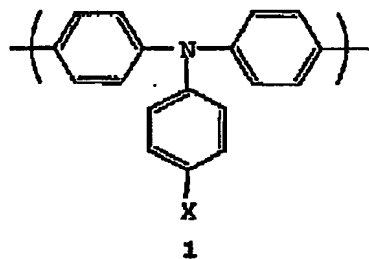
wherein the reaction mixture comprises a catalytic amount of a catalyst suitable for catalysing the polymerisation of the aromatic monomers, and a base in an amount sufficient to convert the boron derivative functional groups into boronate anionic groups.

The inventors have found that polymers according to the invention function effectively as electron transporting materials for red, green or blue electroluminescent polymers without adversely affecting device properties as has been found for systems having aliphatic electron withdrawing 9-substituents such as perfluoroalkyl.

Detailed Description of the Invention

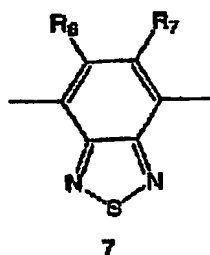
The polymers prepared using monomers according to the invention may be homopolymers or copolymers. A wide range of co-monomers for polymerisation with the monomers of the invention will be apparent to the skilled person. Examples of comonomers include triarylaminines as disclosed in, for example, WO 99/54385 and heteroaryl units as disclosed in, for example, WO 00/46321 and WO 00/55927.

Particularly preferred triarylamine repeat units for such copolymers include units of formulae 1-6:

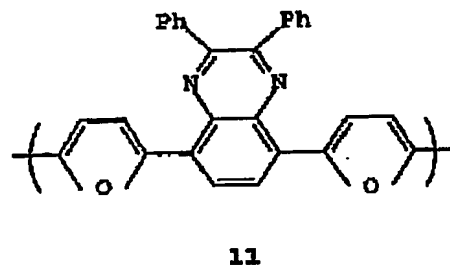
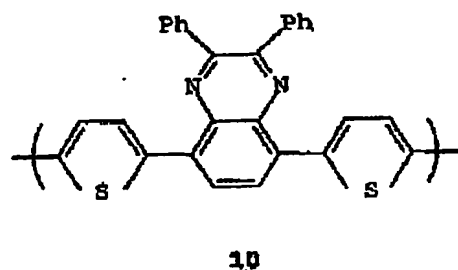
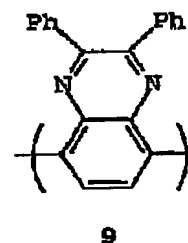
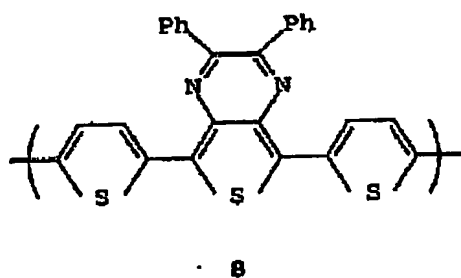


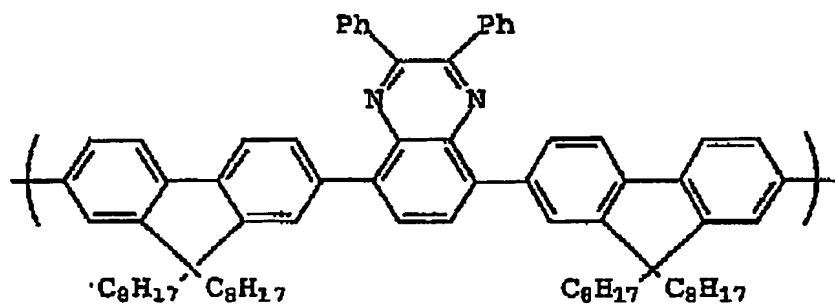
X and Y may be the same or different and are substituent groups. A, B, C and D may be the same or different and are substituent groups. It is preferred that one or more of X, Y, A, B, C and D is independently selected from the group consisting of alkyl, aryl, perfluoroalkyl, thioalkyl, cyano, alkoxy, heteroaryl, alkylaryl and arylalkyl groups. One or more of X, Y, A, B, C and D also may be hydrogen. It is preferred that one or more of X, Y, A, B, C and D is independently an unsubstituted, isobutyl group, an n-alkyl, an n-alkoxy or a trifluoromethyl group because they are suitable for helping to select the HOMO level and/or for improving solubility of the polymer.

Particularly preferred heteroaryl repeat units for such copolymers include units of formulae 7-21:

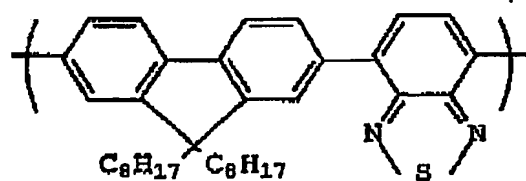


wherein R_8 and R_7 are the same or different and are each independently a substituent group. Preferably, one or both of R_8 and R_7 may be selected from hydrogen, alkyl, aryl, perfluoroalkyl, thioalkyl, cyano, alkoxy, heteroaryl, alkylaryl, or arylalkyl. These groups are preferred for the same reasons as discussed in relation to X, Y, A, B, C and D above. Preferably, for ease of manufacture, R_8 and R_7 are the same. More preferably, they are the same and are each a phenyl group.

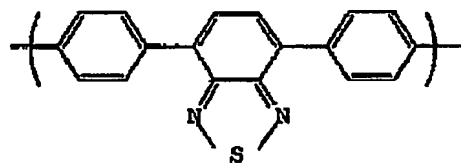




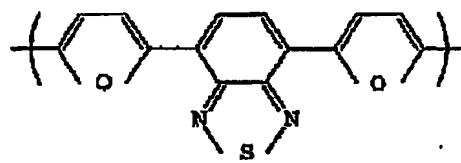
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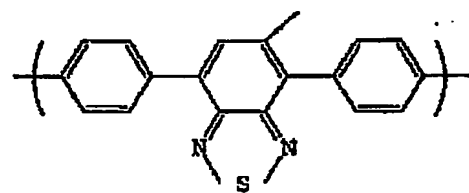
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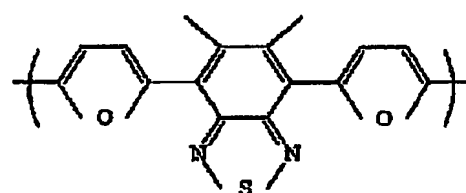
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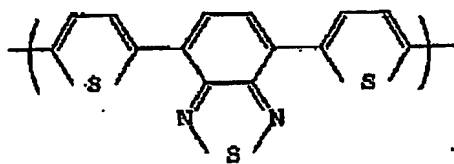
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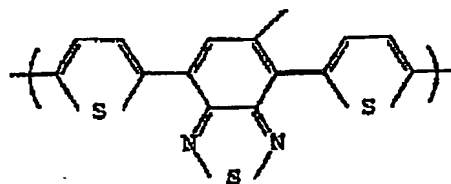
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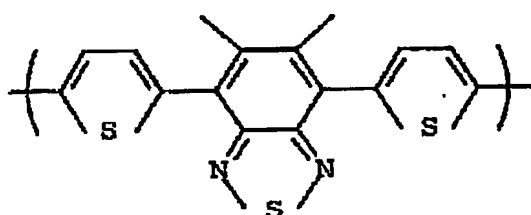
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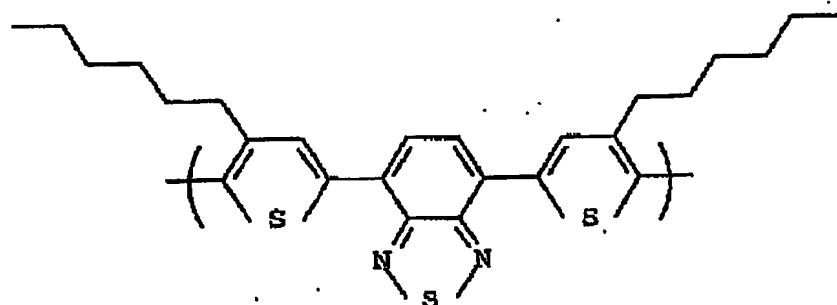
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21

Electron withdrawing groups / heteroaryls suitable for monomers and repeat units of formula (I) according to the invention will be apparent to the skilled person. In particular, those substituents / heteroaryls having a positive Hammett sigma constant may be suitable. The electron withdrawing groups / heteroaryls should preferably be selected to avoid interference with the polymerization of the monomer, e.g. by steric hindrance.

Electron withdrawing groups Ar according to (a) or (b) may be provided with solubilising groups. Particularly preferred as solubilising groups are optionally substituted, branched or linear C₁₋₂₀ alkyl or alkoxy, more preferably C₄₋₁₀ alkyl.

The polymer according to the invention may be a homopolymer or a copolymer. Where it is a copolymer, it may be a 1:1 copolymer, random or block copolymer. A block copolymer according to the invention may comprise at least two regions selected from:

- a hole transporting region
- an electron transporting region
- an emissive region.

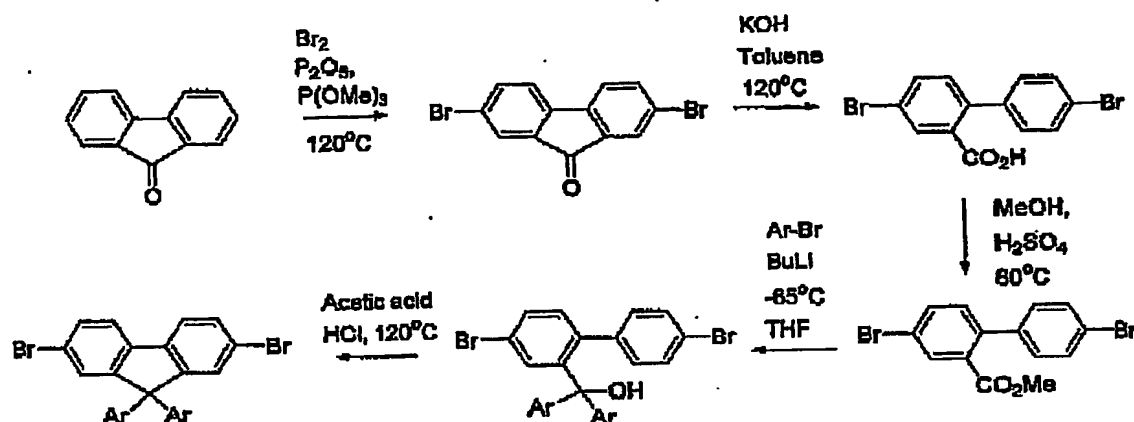
The functions of charge transport and emission may be provided by a range of moieties which will be apparent to the skilled person, as described in, for example, WO 00/55927 or US 6353083.

Preferred methods for polymerisation of the monomers according to the invention are Suzuki polymerisation as described in, for example, WO 00/53656 and Yamamoto polymerisation as described in, for example, T. Yamamoto, "Electrically Conducting And Thermally Stable π -Conjugated Poly(arylene)s Prepared by Organometallic Processes", Progress in Polymer Science 1993, 17, 1153-1205.

Examples

Monomer examples

Monomers according to the invention may be prepared in accordance with the following scheme:



Modelled examples

The effect of appending various groups to the phenyl rings of 9,9-diphenylfluorene repeat unit HOMO and LUMO levels was calculated using AM1 from the AMPAC software package (1) and ZINDO calculations from the Gaussian software package (2).

1) AM1 in Ampac program package

Ampac 5.0 User's Manual, © 1994 Semichem, 7128 Summit, Shawnee, KS 66216

2) ZINDO from Gaussian software:

Gaussian 98, Revision A.9,

M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria,

M. A. Robb, J. R. Cheeseman, V. G. Zakrzewski, J. A. Montgomery, Jr.,

R. E. Stratmann, J. C. Burant, S. Dapprich, J. M. Millam,

A. D. Daniels, K. N. Kudin, M. C. Strain, O. Farkas, J. Tomasi,

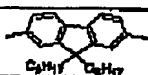
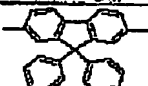


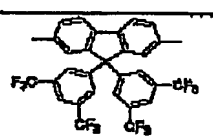
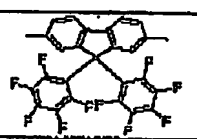
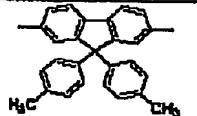
V. Barone, M. Cossi, R. Cammi, B. Mennucci, C. Pomelli, C. Adamo,

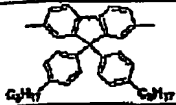
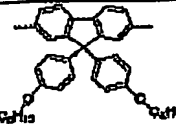
S. Clifford, J. Ochterski, G. A. Petersson, P. Y. Ayala, Q. Cui,

K. Morokuma, D. K. Mallick, A. D. Rabuck, K. Raghavachari,

J. B. Foresman, J. Cioslowski, J. V. Ortiz, A. G. Baboul,
 B. B. Stefanov, G. Liu, A. Liashenko, P. Piskorz, I. Komaromi,
 R. Gomperts, R. L. Martin, D. J. Fox, T. Keith, M. A. Al-Laham,
 C. Y. Peng, A. Nanayakkara, M. Challacombe, P. M. W. Gill,
 B. Johnson, W. Chen, M. W. Wong, J. L. Andres, C. Gonzalez,
 M. Head-Gordon, E. S. Replogle, and J. A. Pople,
 Gaussian, Inc., Pittsburgh PA, 1998.

The results are summarised in the table below:

Structure	Name	HOMO (eV)	LUMO (eV)
	FB	-7.84	0.04
	dpf	-7.79	-0.06
	pdpf	-8.16	-0.40
	mdpf	-8.08	-0.30
	dmdpf	-8.36	-0.72
	fdpf	-8.25	-1.10
	Cl dpf	-7.70	-0.01

	C8dpf	-7.68	0.00
	hadpf	-7.69	0.04

As can be seen from these examples, replacement of octyl with phenyl has a relatively small effect on LUMO level, however a significant change is only effected by substitution by electron withdrawing groups such as fluorine or perfluoroalkyl. By comparison with unsubstituted diphenylfluorene, it can be seen that alkoxy groups substituted in the para position, as per the prior art, are not predicted to show any electron withdrawing character. This is consistent with known electron withdrawing properties of such substituents, in particular their negative Hammett sigma constants.

Polymer Examples

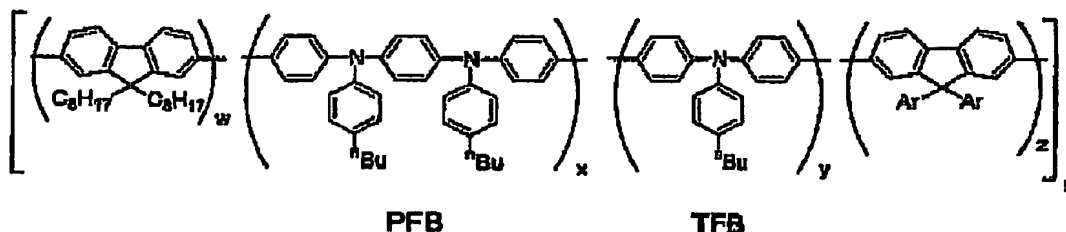
Polymers according to the invention having formula P1 were prepared by Suzuki polymerisation in accordance with the process described in WO 00/53666, by reaction of the following monomers in the ratios set out in the table below:

2,7-dioxalaborane-9,9-di-(n-octyl)fluorene

2,7-dioxalaborane-9,9-di-(4-trifluoromethylphenyl)fluorene

N,N-di(4-bromophenyl)-N-(4-sec-butylphenyl)amine (to produce the "TFB" repeat unit shown below)

Di [N-(4-bromophenyl)-N-(4-n-butylphenyl)]-phenylene-1,4-diamine (to produce the "PFB" repeat unit shown below)

**P1**

wherein $w + x + y + z = 1$, $w + z \geq 0.5$, $0 \leq x + y \leq 0.5$, $z > 0$ and $n \geq 2$

Particular embodiments are as follows:

Example	w	x	y	z
1	50	10	10	30
2	50	0	10	40
3	0	0	50	50
4	0	0	0	100
5	50	0	0	50

Where TFB and PFB are present, as in example 1, the polymer may function as a blue electroluminescent polymer as described in WO 00/55927.

Where TFB is present and PFB is absent the polymer may be a block copolymer with hole and electron transporting segments (example 2) or a 1:1 regioregular hole transporting copolymer (example 3). It may also show blue electroluminescence.

Where TFB and PFB are absent, the polymer may be used as an electron transporting polymer for a red, green or blue electroluminescent material (examples 4 and 5).

Device Example

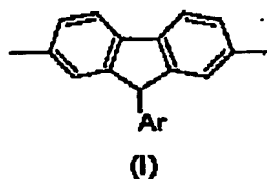
A device according to the invention was prepared as follows:

- 1) Depositing poly(ethylenedioxythiophene) / polystyrene sulfonate (PEDT / PSS), available from Bayer ® as Baytron P ®, by spin coating onto an indium tin oxide anode supported on a glass substrate (available from Applied Films, Colorado, USA).
- 2) Depositing polymer P1 onto the PEDT / PSS by spin coating from xylene solution having a concentration of 2 % w / v.
- 3) Depositing a cathode comprising a first layer of calcium and a second layer of aluminium by evaporation onto the polymer P1.

Although the present invention has been described in terms of specific exemplary embodiments, it will be appreciated that various modifications, alterations and / or combinations of features disclosed herein will be apparent to those skilled in the art without departing from the spirit and scope of the invention as set forth in the following claims.

Claims

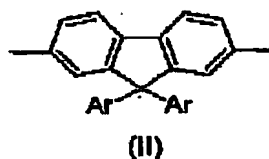
- 1) A polymer comprising optionally substituted first repeat units of formula (I):



wherein Ar is selected from:

- (a) aromatic hydrocarbon substituted with at least one electron withdrawing group or
- (b) electron withdrawing heteroaryl.

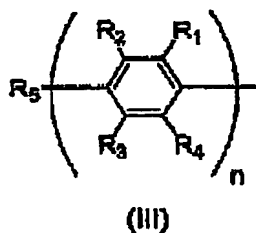
- 2) A polymer according to claim 1 comprising repeat units of formula (II):



wherein each Ar is independently selected from:

- (c) aromatic hydrocarbon substituted with at least one electron withdrawing group or
- (d) electron withdrawing heteroaryl.

- 3) A polymer according to claim 1 or 2 wherein each Ar is independently selected from units of formula (III):



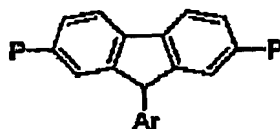
wherein n is from 1-3 and R₁-R₅ are independently selected from:

- hydrogen;
- solubilising groups selected from alkyl, alkoxy, arylalkyl and heteroarylalkyl; and
- electron withdrawing groups

such that at least one of R_1 - R_5 is an electron withdrawing group.

- 4) A polymer according to any preceding claim wherein Ar is phenyl or oligophenyl substituted with at least one electron withdrawing group and the at least one electron withdrawing group is selected from: groups comprising fluorine, cyano and nitro.
- 5) A polymer according to claim 4 wherein the at least one electron withdrawing group is selected from fluorine atoms, fluoroalkyl, fluoroaryl and fluoroheteroaryl.
- 6) A polymer according to claim 1 or 2 wherein Ar is an electron withdrawing heteroaryl selected from optionally substituted pyridines and triazines.
- 7) A polymer according to any preceding claim comprising a second repeat unit.
- 8) A polymer according to claim 7 wherein the second repeat unit is selected from triarylamine and heteroaromatics.
- 9) A polymer according to any preceding claim that is capable of transporting electrons.
- 10) A polymer according to claim 9 that comprises at least one segment capable of hole transport and / or emission.
- 11) An optical device comprising a polymer according to any one of claims 1 to 10.
- 12) An optical device according to claim 11 that is an electroluminescent device.
- 13) An electroluminescent device comprising:
 - a first electrode for injecting charge carriers of a first type;
 - a second electrode for injecting charge carriers of a second type; and
 - an emissive layer comprising a polymer according to any one of claims 1-8 between the first and second electrodes.

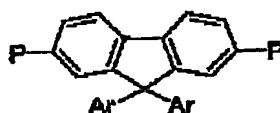
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14) A monomer comprising an optionally substituted compound of formula (IV):



(IV)

wherein each P independently represents a polymerisable group and Ar is as defined in any one of claims 1-8.

15) A monomer according to claim 14 comprising an optionally substituted compound of formula (V):



(V)

wherein each P independently represents a polymerisable group.

16) A monomer according to claim 14 or 15 wherein each P is independently selected from a reactive boron derivative group selected from a boronic acid group, a boronic ester group and a borane group and a reactive halide group.

17) A process for preparing a polymer comprising a step of reacting a first monomer as defined in any one of claims 14-16 with a second monomer that may be the same or different from the first monomer under conditions so as to polymerise the monomers.

18) A process for preparing a polymer according to claim 17 which comprises polymerising in a reaction mixture:

(a) a monomer according to claim 16 wherein each P is a boron derivative functional group selected from a boronic acid group, a boronic ester group and a borane group, and an aromatic monomer having at least two reactive halide functional groups; or

- (b) a monomer according to claim 16 wherein each P is a reactive halide functional group, and an aromatic monomer having at least two boron derivative functional groups selected from boronic acid groups, boronic ester groups and borane groups; or
- (c) a monomer according to claim 16 wherein one P is a reactive halide functional group and one P is a boron derivative functional group selected from a boronic acid group, a boronic ester group and a borane group,

wherein the reaction mixture comprises a catalytic amount of a catalyst suitable for catalysing the polymerisation of the aromatic monomers, and a base in an amount sufficient to convert the boron derivative functional groups into boronate anionic groups.



PCT Application

GB0304753

